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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/551,898	01/05/2006	Kuniharu Moriwaki	10873.1788USWO	6265	
	7590 04/10/200 UMANN, MUELLER	EXAMINER			
P.O. BOX 2902			WILSON, LARRY ROSS		
MINNEAPOLIS, MN 55402-0902			ART UNIT	PAPER NUMBER	
			3767		
			MAIL DATE	DELIVERY MODE	
			04/10/2009	PAPER	

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Office Action Summany		Applicati	on No.	Applicant(s)				
		10/551,8	98	MORIWAKI ET AL.				
Office Action Summary				Art Unit				
			. WILSON	3767				
Period fo	The MAILING DATE of this communication or Reply	n appears on the	e cover sheet with the c	orrespondence ad	ldress			
A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION. - Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication. - Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).								
Status								
1) 🔀	Responsive to communication(s) filed on 3	30 March 2009						
•	. · · · · · · · · · · · · · · · · · · ·							
	/ 							
٥/ك	closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213.							
Disnositi	on of Claims		,,					
•	Claim(s) 1 and 4-7 is/are pending in the application.							
	4a) Of the above claim(s) is/are withdrawn from consideration.							
· · · · · · · · · · · · · · · · · · ·	5) Claim(s) is/are allowed.							
="	Claim(s) <u>1 and 4-7</u> is/are rejected.							
-	Claim(s) is/are objected to.							
8) Claim(s) are subject to restriction and/or election requirement.								
Applicati	on Papers							
9) 🔲 🤈	The specification is objected to by the Exa	miner.						
10)⊠ The drawing(s) filed on <u>20 August 2008</u> is/are: a)⊠ accepted or b)⊡ objected to by the Examiner.								
	Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).							
	Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).							
11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.								
Priority u	ınder 35 U.S.C. § 119							
 12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a) All b) Some * c) None of: 1. Certified copies of the priority documents have been received. 2. Certified copies of the priority documents have been received in Application No 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). 								
* S	See the attached detailed Office action for a	•		d.				
Attachmen	t(s)							
1) Notice of References Cited (PTO-892) 4) Interview Summary (PTO-413)								
3) 🔲 Inforr	e of Draftsperson's Patent Drawing Review (PTO-948 nation Disclosure Statement(s) (PTO/SB/08) r No(s)/Mail Date	3)	Paper No(s)/Mail Da 5) Notice of Informal P 6) Other:					

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DETAILED ACTION

Claim Rejections - 35 USC § 103

1. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all

obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person

such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the

manner in which the invention was made.

2. Claims 1, and 4-7 are rejected under 35 U.S.C. 103(a) as being unpatentable over

European Patent Application EP 1 048 311 A2 to Yosisuke Teraoka (Teraoka) in view of

Whisson.

In regards to claim 1, Teraoka teaches a medical needle device with a winged

shield (Fig. 1, #1) comprising a winged shield (Fig. 1, #7 & 8), that has a substantially

cylindrical shield tube (Fig. 1, #8) an a pair of wings (Fig. 1, #7), a hub that is inserted

into an inner bore of the shield tube so as to be movable in an axial direction (col. 2, lines

22-27), a needle that is mounted to a front end of the hub (Fig. 1, #3), a rear end of the

hub capable of being connected with an infusion tube (col. 7, lines 43-44) and a tip of the

needle capable of being stored in the inner bore of the shield tube (col. 2, lines 26-27), the

needle is inserted into and coupled with a bore of the hub at a front end thereof (Fig. 5,

#11, 12 – shows the needle 11 inserted into the hub 12).

But Teraoka does not teach wherein at least a part of the hub is made of a material having

flexibility, wherein that the shield tube and the hub are bendable together at least in a part

of a range along an axial direction when the needle protrudes from the front end of the

shield tube and is latched to shield tube.

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Whisson teaches wherein at least a part of the hub is made of a material having flexibility (col. 2, line 36), wherein the shield tube and the hub are bendable together at least in a part of a range along an axial direction when the needle protrudes from the front end of the shield tube (Fig. 6 shows the extended position of the needle & col. 3, lines 11-16 — a flexible delivery tube within a flexible tubular duct are bendable) and is latched to the shield tube (col. 4, lines 24-28).

It would have been obvious to one of ordinary skill in the art at the time of the invention to have included the flexible delivery tube and the flexible duct of Whisson as a hub and a shield tube, respectively, in the medical needle of Teraoka in order to allow for destructive bending or "kinking" of the delivery tube to render the infusion set incapable of further use (col. 4, lines 61-64) as explicitly taught by Whisson.

In regards to claims 2-7, Teraoka, as modified by Whisson teaches the medical needle device according to claim 1 (see rejection above), the shield tube is made of material having flexibility (col. 3, lines 11-16 — a flexible delivery tube within a flexible tubular duct are bendable, the flexible tubular duct is a part of the shield tube); Teraoka further teaches wherein the shield tube (Fig. 1, #8) includes an extendable portion that is structured to be extendable and contractible (col. 5, lines 24-27), the needle can be moved in the axial direction of the shield tube by extending and contracting the extendable portion (col. 5, lines 27-29) and the shield tube and the hub are bendable at the extendable portion (implied the shield tube, as modified by Whisson, is flexible, the extendable portion of Teraoka is flexible otherwise it could not be extendable thus both are bendable); wherein the extendable portion has a plasticity-process accordion-like

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structure (col. 5, lines 31-35); and when the shield tube and the hub in the inner bore of the shield tube are bent together, a minimum radius of curvature at a bent part can be 3 mm or smaller (implied in the flexible nature of the fluid delivery tube that holds the needle, as modified by Whisson, and the bendable accordion structure of the extendable member is capable of bending 3 mm or smaller, furthermore it would have been obvious to one of ordinary skill in the art to chose materials that would optimize the curve to allow the patient a greater range of motion such that conscious and unconscious movement does not remove the needle or damaging the vein. See MPEP 2144.05 II A -

3. The amendment of claim 1 in the amendment filed on 30 March 2009 is acknowledged.

Response to Amendment

The amendment of claim 1 and cancellation of claims 2 and 3 in the amendment filed on 8

January 2009 is acknowledged.

Optimization of ranges).

a. Claim Rejections - 35 USC § 102

The amendment to claim 1 overcomes the rejection which is hereby withdrawn.

Response to Arguments

4. Applicant's arguments filed 30 March 2009 have been fully considered but they are not

persuasive. The applicant's argument that there is no reasonable basis to modify the older and

tube in Teraoka with the flexible delivery tube and duct of Whisson is not persuasive because

Whisson provides a reason for the flexible delivery tube such as to render the infusion set

incapable of further use by kinking the delivery tube, which is in addition to the needle retention

elements in the base 11. Additionally, if the flexible delivery tube and flexible delivery duct of

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Whisson would not provide adequate support when used in combination with the device of

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Teraoka it would not provide adequate support of the device according to Whisson either,

because it would continue to function in the same way as it does in Whisson.

Conclusion

5. Any inquiry concerning this communication or earlier communications from the

examiner should be directed to LARRY R. WILSON whose telephone number is (571)270-5899.

The examiner can normally be reached on Monday-Thursday 7:00 AM - 5:30 PM (EST).

6. If attempts to reach the examiner by telephone are unsuccessful, the examiner's

supervisor, Kevin C. Sirmons can be reached on 571-272-4965. The fax phone number for the

organization where this application or proceeding is assigned is 571-273-8300.

7. Information regarding the status of an application may be obtained from the Patent

Application Information Retrieval (PAIR) system. Status information for published applications

may be obtained from either Private PAIR or Public PAIR. Status information for unpublished

applications is available through Private PAIR only. For more information about the PAIR

system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR

system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would

like assistance from a USPTO Customer Service Representative or access to the automated

information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/LARRY R WILSON/ Examiner, Art Unit 3767

Examiner, Art Omt 3707

/Kevin C. Sirmons/

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Supervisory Patent Examiner, Art Unit 3767

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